

1 **Adaptation of Host Transmission Cycle During *Clostridium difficile***

2 **Speciation**

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43 Bacterial speciation is a fundamental evolutionary process characterized by diverging
44 genotypic and phenotypic properties. However, the selective forces impacting genetic
45 adaptations and how they relate to the biological changes underpinning the formation of a
46 new bacterial species remain poorly understood. Here we show that the spore-forming,
47 healthcare-associated enteropathogen *Clostridium difficile* is actively undergoing speciation.
48 Applying large-scale genomic analysis of 906 strains, we demonstrate that the ongoing
49 speciation process is linked to positive selection on core genes in the newly forming species
50 that are involved in sporulation and the metabolism of simple dietary sugars. Functional
51 validation demonstrates the new *C. difficile* produce more resistant spores and show
52 increased sporulation and host colonization capacity when glucose or fructose is available for
53 metabolism. Thus, we report the formation of an emerging *C. difficile* species, selected for
54 metabolizing simple dietary sugars and producing high levels of resistant spores that is
55 adapted for healthcare-mediated transmission.

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67 The formation of a new bacterial species from its ancestor is characterized by genetic
68 diversification and biological adaptation¹⁻⁴. For decades, a polyphasic examination⁵, relying
69 on genotypic and phenotypic properties of a bacterium, has been used to define and
70 discriminate a “species”. The bacterial taxonomic classification framework has more recently
71 used large-scale genome analysis to incorporate aspects of a bacterium’s natural history, such
72 as ecology⁶, horizontal gene transfer¹, recombination² and phylogeny³. Although a more
73 accurate definition of a bacterial species can be achieved with whole-genome-based
74 approaches, we still lack a fundamental understanding of how selective forces impact
75 adaptation of biological pathways and phenotypic changes leading to bacterial speciation. In
76 this work, we describe the genome evolution and biological changes during the ongoing
77 formation of a new *C. difficile* species that is highly specialized for human transmission in the
78 modern healthcare system.

79 *C. difficile* is a strictly anaerobic, Gram-positive bacterial species that produces highly
80 resistant, metabolically dormant spores capable of rapid transmission between mammalian
81 hosts through environmental reservoirs⁷. Over the past four decades, *C. difficile* has emerged
82 as the leading cause of antibiotic-associated diarrhea worldwide, with a large burden on the
83 healthcare system^{7,8}. To define the evolutionary history and genetic changes underpinning the
84 emergence of *C. difficile* as a healthcare pathogen, we performed whole-genome sequence
85 analysis of 906 strains isolated from humans (n = 761), animals (n = 116) and environmental
86 sources (n = 29) with representatives from 33 countries and the largest proportion originating
87 from the UK (n = 465) (Supplementary Fig. 1; Supplementary Table 1; Supplementary Table
88 2). This dataset is summarized visually here <https://microreact.org/project/H1QidSp14>. Our
89 collection was designed to capture comprehensive *C. difficile* genetic diversity⁹ and includes
90 13 high-quality and well-annotated reference genomes (Supplementary Table 2). Robust
91 maximum likelihood phylogeny based on 1,322 concatenated single-copy core genes (Fig.

1a; Supplementary Table 3) illustrates the existence of four major phylogenetic groups within this collection. Bayesian analysis of population structure (BAPS) using concatenated alignment of 1,322 single-copy core genes corroborated the presence of the four distinct phylogenetic groupings (PGs 1-4) (Fig. 1a) that each harbor strains from different geographical locations, hosts and environmental sources which indicates signals of sympatric speciation. Each phylogenetic group also harbors distinct clinically relevant ribotypes (RT): PG1 (RT001, 002, 014); PG2 (RT027 and 244); PG3 (RT023 and 017); PG4 (RT078, 045 and 033).

The phylogeny was rooted using closely related species (*C. bartlettii*, *C. hiranonis*, *C. ghonii* and *C. sordellii*) as outgroups (Fig. 1a). This analysis indicated that three phylogenetic groups (PG1, 2 and 3) of *C. difficile* descended from the most diverse phylogenetic group (PG4). This was also supported by the frequency of single-nucleotide polymorphism (SNP) differences in pairwise comparisons between strains of PG4 and each of the other PGs versus the level of pairwise SNP differences between comparisons of PGs 1, 2 and 3 to each other (Supplementary Fig. 2). Interestingly, bacteria from PG4 display distinct colony morphologies compared to bacteria from PG 1, 2 and 3 when grown on nutrient agar plates (Supplementary Fig. 3), suggesting a link between *C. difficile* colony phenotype and genotype that distinguishes PG 1, 2 and 3 from PG4.

Our previous genomic study using 30 *C. difficile* genomes indicated an ancient, genetically diverse species that likely emerged 1 to 85 million years ago¹⁰. Testing this estimate using our larger dataset indicated the species emerged approximately 13.5 million years (12.7-14.3 million) ago. Using the same BEAST¹¹ analysis on our substantially expanded collection, we estimate the most recent common ancestor (MRCA) of PG4 (using RT078 lineage) arose approximately 385,000 (297,137-582,886) years ago. In contrast, the MRCA of the PG1, 2 and 3 groups (using RT027 lineage) arose approximately 76,000

(40,220-214,555) years ago. Bayesian skyline analysis reveals a population expansion of PG1, 2 and 3 groups (using RT027 lineage) around 1595 A.D., which occurred shortly before the emergence of the modern healthcare system in the 18th century (Supplementary Fig. 4). Combined, these observations suggest that PG4 emerged prior to the other PGs and that the PG1, 2 and 3 population structure started to expand just prior to the implementation of the modern healthcare system¹². We therefore refer to PG1, 2 and 3 groups as *C. difficile* “clade A” and PG4 as *C. difficile* “clade B”.

To investigate genomic relatedness, we next performed pairwise Average Nucleotide Identity (ANI) analysis (Fig. 1b). This analysis revealed high nucleotide identity (ANI > 95%) between PGs 1, 2 and 3 indicating that bacteria from these groups belong to the same species; however, ANI between PG4 and any other PG was either less than the species threshold (ANI > 95%) or on the borderline of the species threshold (94.04%-96.25%) (Fig. 1b). To detect recombination events, FastGEAR analysis¹³ was performed on whole-genome sequences of 906 strains. While analysis identified increased recombination within *C. difficile* clade A (PG1-PG2: 1-102, PG1-PG3: 1-214, PG2-PG3: 1-96) (Supplementary Fig. 5) a restricted number of recombination events between *C. difficile* clade A and clade B was observed (PG1-PG4: 1-20, PG2-PG4: 1-25, PG3-PG4: 1-46). This analysis strongly indicates the presence of recombination barriers in the core genome that further distinguishes the two *C. difficile* clades and could encourage sympatric speciation. Furthermore, accessory genome functional analysis also shows a clear separation between clade A and clade B (Supplementary Fig. 6; Supplementary Table 4-5). We also observe a higher number of pseudogenes in clade A compared to clade B (Supplementary Fig. 7; Supplementary Table 6-11). Taken together, these results indicate different selection pressures on the genomes of *C. difficile* clades A and B.

In addition to reduced rates of recombination events, advantageous genetic variants in a population driven by positive selective pressures, termed positive selection, are also a marker of speciation⁶. We determined the Ka/Ks ratios and identified 172 core genes in clade A and 93 core genes in clade B that were positively selected (Ka/Ks >1) (Fig. 2a; Supplementary Table 12-13). Functional annotation and enrichment analysis identified positively selected genes involved in carbohydrate and amino acid metabolism, sugar phosphotransferase system (PTS) and spore coat architecture and spore assembly in clade A (Fig. 2b). In contrast, the sulphur relay system was the only enriched functional category in positively selected genes from clade B. Notably, 26% (45 in total) of the positively selected genes in *C. difficile* clade A produce proteins that are either directly involved in spore production, are present in the mature spore proteome¹⁴ or are regulated by Spo0A¹⁵ or its sporulation-specific sigma factors¹⁶ (Fig. 2c). In contrast, no positively selected genes are directly involved in spore production in *C. difficile* clade B; however, 22.5% (21 genes in total) are either present in the mature spore proteome or are regulated by Spo0A or its sporulation specific sigma factors (Supplementary Fig. 8). The lack of overlap between sporulation-associated positively selected genes in the two lineages suggests a divergence of spore-mediated transmission functions. In addition, these results suggest functions important for host-to-host transmission have evolved in *C. difficile* clade A.

We found 20 positively selected genes (Supplementary Table 12) in clade A whose products are components of the mature spore^{14,15} and could contribute to environmental survival¹⁷. As an example, *sodA* (superoxide dismutase A), a gene associated with spore coat assembly, has three-point mutations which are present in all clade A genomes but absent in clade B genomes (Supplementary Fig. 9). Spores derived from diverse *C. difficile* clades have a wide variation in resistance to microbiocidal free radicals from gas plasma¹⁸. To investigate if the phenotypic resistance properties of spores from the new lineage have evolved, we

exposed spores from both clades to hydrogen peroxide, a commonly used healthcare environmental disinfectant¹⁷. Spores derived from clade A were more resistant to 3% ($P = 0.0317$) and 10% hydrogen peroxide ($P = 0.0317$) when compared to spores from clade B, although there was no difference in survival at 30% peroxide likely due to the overpowering bactericidal effect at this concentration ($P = 0.1667$) (Fig. 3a).

The master regulator of *C. difficile* sporulation, *Spo0A*, is under positive selection in *C. difficile* clade A only. *Spo0A* also controls other host colonization factors, such as flagella, and carbohydrate metabolism, potentially serving to mediate cellular processes to direct energy to spore production and host colonization to facilitate host-to-host transmission¹⁵. Interestingly, the clade A genomes contain genes under positive selection that are involved in fructose metabolism (*fruABC* and *fruK*), glycolysis (*pgk* and *pyk*), sorbitol (CD630_24170) and ribulose metabolism (*repI*), and conversion of pyruvate to lactate (*ldh*). To further explore the link between sporulation and carbohydrate metabolism in clade A, we analyzed positively selected genes using KEGG pathways¹⁹ and manual curation. Manual curation of key enriched pathways across the 172 positively selected core genes in *C. difficile* clade A identified a complete fructose-specific PTS pathway and identified four genes (30%, 4/13) involved in anaerobic glycolysis during glucose metabolism (Supplementary Fig. 10). Other genes associated with enriched PTS pathways include genes used for the cellular uptake and metabolism of mannitol, cellobiose, glucitol/sorbitol, galactitol, mannose and ascorbate. Furthermore, comparative analysis of carbohydrate active enzymes (CAZymes)²⁰ identified a clear separation of CAZymes between *C. difficile* clade A and clade B (Supplementary Fig. 11; Supplementary Table 14). Combined, these observations suggest a divergence of functions between two *C. difficile* clades linked to metabolism of a broad range of simple dietary sugars²¹.

190 The simple sugars glucose and fructose are increasingly used in diets within Western
191 societies²¹. Interestingly, trehalose, a disaccharide of glucose, used as a food additive has
192 impacted the emergence of some human virulent *C. difficile* variants²². Based on our genomic
193 analysis, we hypothesized that dietary glucose or fructose could differentially impact host
194 colonization by spores from *C. difficile* clade A or clade B. We therefore supplemented the
195 drinking water of mice with either glucose, fructose or ribose and challenged with clade A or
196 clade B strains. Ribose metabolic genes were not under positive selection so this sugar was
197 included as a control. Mice challenged with clade A spores exhibited increased bacterial load
198 when exposed to dietary glucose ($P = 0.048$) or fructose ($P = 0.0045$) compared to clade B
199 (Fig. 3b). No difference in bacterial load was observed between *C. difficile* clade A and clade
200 B without supplemented sugars or when supplemented with ribose ($P = 0.2709$) (Fig. 3b).

201 The infectivity and transmission of *C. difficile* within healthcare settings is facilitated
202 by environmental spore density^{23,24}. To determine the impact of simple sugar availability on
203 spore production rates we assessed the ability of the two lineages to form spores in basal
204 defined medium (BDM) alone or supplemented with either glucose, fructose or ribose. While
205 no difference was observed on the ribose control ($P = 0.3095$), *C. difficile* clade A strains
206 exhibited increased spore production on glucose ($P = 0.0317$) or fructose ($P = 0.0317$) (Fig.
207 3c). These results provide experimental validation and, together with our genomic
208 predictions, suggest that enhanced host colonization and onward spore-mediated transmission
209 with the consumption of simple dietary sugars is a feature of *C. difficile* clade A but not clade
210 B.

211 The rapid recent emergence of *C. difficile* as a significant healthcare pathogen has
212 mainly been attributed to the genomic acquisition of antibiotic resistance and carbohydrate
213 metabolic functions on mobile elements via horizontal gene transfer^{22,25}. Here we show that
214 these recent genomic adaptations have occurred in established, distinct evolutionary lineages

each with core genomes expressing unique, pre-existing transmission properties. We reveal the ongoing formation of a new species with biological and phenotypic properties consistent with a transmission cycle that was primed for human transmission in the modern healthcare system (Fig. 3d). Indeed, different transmission dynamics and host epidemiology have also been reported for *C. difficile* clade A (027 lineage²⁶ and 017 lineage²⁷) endemic in healthcare systems in different parts of the world, and the 078 lineage that likely enters the human population from livestock²⁸⁻³⁰. Further, broad epidemiological screens of *C. difficile* present in the healthcare system often highlight high abundances of *C. difficile* clade A as they represent 68.5% (USA), 74% (Europe) and 100% (Mainland China) of the infecting strains^{7,8,31,32}. Thus, we report a link between *C. difficile* clade A speciation, adapted biological pathways and epidemiological patterns. In summary, our study elucidates how bacterial speciation may prime lineages to emerge and transmit in a process accelerated by modern human diet, the acquisition of antibiotic resistance or healthcare regimes.

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254 **Author Contributions**

255 N.K. and T.D.L. conceived and managed the study. N.K., S.C.F., E.V., H.P.B. and T.D.L.
256 wrote the manuscript. D.J.F., P.R., M.P., M.R.J.C., M.B.F.J., K.R.H., M.I., L.H.W., C.S.,
257 T.N., G.D., T.V.R., E.J.K., B.W.W. provided critical input and contributed to the editing of
258 the manuscript. N.K. performed the computational analysis. H.P.B. performed genome
259 annotation of reference genomes. D.J.F., P.R., M.P., M.R.J.C., M.B.F.J., K.R.H., M.I.,
260 L.H.W., C.S., T.N. provided *C. difficile* strains. E.V., H.P.B., S.C.F. and T.D.L. designed *in*
261 *vitro* and *in vivo* experiments. H.P.B., E.V. and M.S. performed *in vitro* experiments. E.V.,
262 M.D.S., S.C. and K.H. performed *in vivo* experiments.

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264 **Conflict of interests**

The authors declare no competing financial interests.

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Figure legends:

Figure 1. Phylogeny and population structure of *Clostridium difficile*. (a) Maximum likelihood tree of 906 *C. difficile* strains constructed from the core genome alignment, excluding recombination events. Collapsed clades as triangles represent four Phylogenetic groups (PG1-4) identified by Bayesian analysis of population structure (BAPS). Number in parentheses indicates number of strains. Key PCR ribotypes in each PG are shown. Bootstrap values of key branches are shown next to the branches. Dates indicate estimated emergence of *C. difficile* species-13.5 million (range 12.7-14.3) years ago, PG4- 385,000 (range 297,137-582,886) years ago and PG1-3- 76,000 (range 40,220-214,555) years ago. *C. bartlettii*, *C. hiranonis*, *C. ghonii* and *C. sordellii* were used as outgroups to root the tree. Scale bar indicates number of substitutions per site. (b) Distribution pattern of average nucleotide identity (ANI) for 906 *C. difficile* strains. Pairwise ANI calculations between different PGs are shown. Dotted red line indicates bacterial species cut-off.

Figure 2. Adaptation of sporulation and metabolic genes in *Clostridium difficile* clade A. Positive selection analysis of *C. difficile* clade A and B based on 1,322 core genes. (a) Distribution of Ka/Ks ratio for the positively selected genes in *C. difficile* clade A (n = 172 genes) and clade B (n = 93 genes) is shown. Error bars are standard error of the mean (SEM).

(b) Enriched functions in the positively selected genes of *C. difficile* clade A (n = 172 genes) and clade B (n = 93 genes) are shown. Y-axis represents number of positive selected genes in each enriched function. All are statistically significant (sugar phosphotransferase system (q = 0.00167), fructose and mannose metabolism (q = 0.001173), sporulation, differentiation and germination (q = 0.0165), cysteine and methionine metabolism (q = 0.00279), sulphur relay system (q = 0.00791)). One-sided Fisher's exact test with *P* value adjusted by Hochberg method. (c) Positively selected sporulation-associated genes in *C. difficile* clade A are shown in blue. Of the 172 genes under positive selection, 26% (45 in total) are either involved in spore production (sporulation stages I, III, IV and V), their proteins are present in the mature spore proteome or they are regulated by Spo0A or its sporulation specific sigma factors.

Figure 3. Bacterial speciation is linked to increased host adaptation and transmission

ability. (a) Spores of *C. difficile* clade A are more resistant to widely used hydrogen peroxide disinfectant. Spores of *C. difficile* clade A and clade B (n = 5 different ribotypes for both lineages) were exposed to hydrogen peroxide for 5 minutes, washed and plated. Recovered CFUs representing surviving germinated spores were counted and presented as a percentage of spores exposed to PBS. Mean and range of 3 independent experiments is presented, Mann-Whitney unpaired two-tailed test. (b) Intestinal colonization of clade A strains is increased in the presence of simple sugars compared to clade B strains. Comparison of vegetative cell loads between *C. difficile* clade A (n = 1, RT027) and clade B (n = 1, RT078) strains in mice whose diet was supplemented with different sugars before challenging with spores. CFUs from fecal samples cultured 16 hours after *C. difficile* challenge are presented. Mean values of 5 mice are presented from 1 representative experiment which was repeated once with similar results, standard error of the mean (SEM), unpaired two-tailed *t* test. (c) Clade A strains produce more spores in the presence of simple sugars. *C. difficile* clade A and clade B

(n = 5 different ribotypes for both lineages) strains were grown on basal defined media in the presence or absence of different sugars, vegetative cells were killed by ethanol exposure and the number of CFUs representing germinated spores were counted. The percentage of spores recovered in the presence of sugars compared to BDM alone is presented. Mean and range of 3 independent experiments is presented, Mann-Whitney unpaired two-tailed test. (d) Overview of adaptations in key aspects of the *C. difficile* clade A transmission cycle in human population.

Online Methods

Collection of *C. difficile* strains

Laboratories worldwide were asked to send a diverse representation of their *C. difficile* collections to the Wellcome Sanger Institute (WSI). After receiving all shipped samples the DNA extraction was performed batch-wise using the same protocol and reagents to minimize bias. Phenol-Chloroform was the preferred method for extraction since it provides high DNA yield and intact chromosomal DNA.

The genomes of 382 strains designated as *C. difficile*, by PCR ribotyping were sequenced and combined with our previous collection of 506 *C. difficile* strains, 13 high quality *C. difficile* reference strains and 5 publicly available *C. difficile* RT 244 strains making a total of 906 strains analyzed in this study. This genome collection includes strains from humans (n = 761), animals (n = 116) and the environment (n = 29) that were collected from diverse geographic locations (UK; n = 465, Europe; n = 230, N-America; n = 111, Australia; n = 62, Asia; n = 38). Details of all strains are listed in Supplementary Table 1 and Supplementary Table 2, including the European Nucleotide Archive (ENA) sample accession numbers. Metadata of this *C. difficile* collection have been made freely publicly available through Microreact³³ (<https://microreact.org/project/H1QidSp14>).

Bacterial culture and genomic DNA preparation

C. difficile strains were cultured on blood agar plates for 48 hours, inoculated into brain–heart infusion broth supplemented with yeast extract and cysteine and grown overnight (16 hours) anaerobically at 37 °C. Cells were pelleted, washed with PBS, and genomic DNA preparation was performed using a phenol–chloroform extraction as previously described³⁴. All culturing of *C. difficile* took place in anaerobic conditions (10% CO₂, 10% H₂, 80% N₂) in a Whitley DG250 workstation at 37 °C. All reagents and media were reduced for 24 hours in anaerobic conditions before use.

DNA sequencing, assembly and annotation

Paired-end multiplex libraries were prepared and sequenced using Illumina Hi-Seq platform with fragment size of 200-300 bp and a read length of 100 bp, as previously described^{35,36}. An in-house pipeline developed at the WSI (<https://github.com/sanger-pathogens/Bio-AutomatedAnnotation>) was used for bacterial assembly and annotation. It consisted of *de novo* assembly for each sequenced genome using Velvet v1.2.10³⁷, SSPACE v2.0³⁸ and GapFiller v1.1³⁹ followed by annotation using Prokka v1.5-1⁴⁰. For the 13 high-quality reference genomes, strains Liv024, TL178, TL176, TL174, CD305 and Liv022 were sequenced using 454 and Illumina sequencing platforms, BI-9 and M68 were sequenced using 454 and capillary sequencing technologies with the assembled data for these 8 strains been improved to an ‘Improved High Quality Draft’ genome standard⁴¹. Optical maps using the Argus Optical Mapping system were also generated for Liv024, TL178, TL176, TL174, CD305 and Liv022. The remaining strains are all contiguous and were all sequenced using 454 and capillary sequencing technologies except for R20291 which also had Illumina data incorporated and 630 which was sequenced using capillary sequence data alone.

Phylogenetic analysis, Pairwise SNP distances analysis and Average Nucleotide Identity analysis

The phylogenetic analysis was conducted by extracting nucleotide sequence of 1,322 single copy core gene from each *C. difficile* genome using Roary⁴². The nucleotide sequences were concatenated and aligned with MAFFT v7.20⁴³. Gubbins⁴⁴ was used to mask recombination from concatenated alignment of these core genes and a maximum-likelihood tree was constructed using RAxML v8.2.8⁴⁵ with the best-fit model of nucleotide substitution (GTRGAMMA) calculated from ModelTest embedded in TOPALi v2.5⁴⁶ and 500 bootstrap replicates. The phylogeny was rooted using a distance-based tree generated using Mash v2.0⁴⁷, R package APE⁴⁸ and genome assemblies of closely related species (*C. bartlettii*, *C. hiranonis*, *C. ghonii* and *C. sordellii*). All phylogenetic trees were visualized in iTOL⁴⁹. Genomes of closely related *C. difficile* were downloaded from NCBI. Pairwise SNP distances analysis was performed on concatenated alignment of 1,322 single-copy core genes using SNP-Dist (<https://github.com/tseemann/snp-dists>). Average nucleotide analysis (ANI) was calculated by performing pairwise comparison of genome assemblies using MUMmer⁵⁰.

Population structure and recombination analysis

Population structure based on concatenated alignment of 1,322 single-copy core genes of *C. difficile* was inferred using the HierBAPS⁵¹ with one clustering layers and 5, 10 and 20 expected numbers of clusters (k) as input parameters. Recombination events across the whole-genome sequences were detected by mapping genomes against a reference genome (NCTC 13366; RT027) and using FastGear¹³ with default parameters.

Functional genomic analysis

To explore accessory genome and identify protein domains in a genome, we performed RPS-BLAST using COG database (accessed February 2019)⁵². All protein domains were classified in different functional categories using the COG database⁵² and were used to perform Discriminant Analysis of Principle Components (DAPC)⁵³ implemented in

the R package Adegnet v2.0.1⁵⁴. Domain and functional enrichment analysis was calculated using one-sided Fisher's exact test with *P* value adjusted by Hochberg method in R v3.2.2.

Carbohydrate active enzymes (CAZymes) in a genome were identified using dbCAN v5.0⁵⁵ (HMM database of carbohydrate active enzyme annotation). Best hits include hits with E-value $< 1 \times 10^{-5}$ if alignment > 80 aa and hits with E-value $< 1 \times 10^{-3}$ if alignment < 80 aa, and alignment coverage > 0.3 . Best hits were used to perform Discriminant Analysis of Principle Components (DAPC)⁵³ implemented in the R package Adegnet v2.0.1⁵⁴.

Functional annotation of positively selected genes was carried out using the Riley classification system⁵⁶, KEGG Orthology⁵⁷ and Pfam functional families⁵⁸.

Analysis of selective pressures.

The aligned nucleotide sequences of each 1,322 single copy core genes were extracted from Roary's output. The ratio between the number of non-synonymous mutations (Ka) and the number of synonymous mutations (Ks) was calculated for the whole alignment and for the respective subsets of strains belonging to the PG1, 2, 3 as a group and PG4. The Ka/Ks ratio for each gene alignment was calculated with SeqinR v3.1. A Ka/Ks > 1 was considered as the threshold for identifying genes under positive selection.

Pseudogenes analysis

Nucleotide annotations of genes within a genome within each phylogenetic group were mapped against the protein sequences of the reference genome for its phylogenetic group (PG1: NCTC 13307(RT012), PG2: SRR2751302 (RT244), PG3: NCTC 14169 (RT017), PG4: NCTC 14173 (RT078)) using TBLASTN as previously described⁵⁹. Pseudogenes were called based on following criteria: genes with E value $> 1 \times 10^{-30}$ and sequence identity $< 99\%$ and which are absent in 90% members of a phylogenetics group. Genes in the reference genomes annotated as a pseudogene were also included in addition to genes in query genomes.

Analysis of estimating dates

The aligned nucleotide sequences of each 222 core genes of *C. difficile* which are under neutral selection ($Ka/Ks = 1$) were extracted from Roary's output. Gubbins⁴⁴ was used to mask recombination from concatenated alignment of these core genes and used as an input for Bayesian Evolutionary Analysis Sampling Trees (BEAST) software package v2.4.1¹¹. In BEAST, the MCMC chain was run for 50 million generations, sampling every 1,000 states using the strict clock model ($2.50 \times 10^{-9} - 1.50 \times 10^{-8}$ per site per year)¹⁰ and HKY four discrete gamma substitution model, each run in triplicate. Convergence of parameters were verified with Tracer v1.5⁶⁰ by inspecting the Effective Sample Sizes ($ESS > 200$). LogCombiner was used to remove 10% of the MCMC steps discarded as burn-ins and combine triplicates. The resulting file was used to infer the time of divergence from the most recent common ancestor for *C. difficile*, *C. difficile* clade A and clade B. The Bayesian skyline plot was generated with Tracer v1.5⁶⁰.

***C. difficile* growth in vitro on selected carbon sources**

Basal defined medium (BDM)⁶¹ was used as the minimal medium to which selected carbon sources (2 g/l of glucose, fructose or ribose from Sigma-Aldrich) were added. *C. difficile* strains were grown on CCEY agar (Bioconnections) for two days; 125-ml Erlenmeyer flasks containing 10 ml of BDM with or without carbon sources were inoculated with *C. difficile* strains and incubated in anaerobic conditions at 37 °C shaking at 180 rpm. After 48 hours, spores were enumerated by centrifuging the culture to a pellet, carefully decanting the BDM and re-suspending in 70% ethanol for 4 hours to kill vegetative cells. Following ethanol shock, spores were washed twice in PBS and plated in a serial dilution on YCFA media⁶² supplemented with 0.1% sodium taurocholate. Colony forming units (representing germinated spores) were counted 24 hours later. The experiment was performed independently 3 times for each strain. Clade A strains used were TL178 (RT002/ PG1),

TL174 (RT015/ PG1), R20291 (RT027/ PG2), CF5 (RT017/ PG3) and CD305 (RT023/ PG3). Clade B strains used were MON024 (RT033), CDM120 (RT078), WA12 (RT291), WA13 (RT228) and MON013 (RT127). Data were presented using GraphPad Prism v7.03.

***C. difficile* spore resistance to disinfectant**

Spores were prepared by adapting the previous protocol¹⁸. In brief, *C. difficile* strains were streaked on CCEY media, the cells were harvested from the plates 48 hours later and subjected to exposure in 70% ethanol for 4 hours to kill vegetative cells. The solution was then centrifuged, ethanol was decanted and the spores were washed once in 5 ml sterile saline (0.9% w/v) solution before being suspended in 5 ml of saline (0.9% w/v) with Tween20 (0.05% v/v). 300 µl spore suspensions (at a concentration of approximately 10⁶ spores) were exposed to 300 µl of 3%, 10% and 30% hydrogen peroxide (Fisher Scientific UK Limited) solutions for 5 minutes in addition to 300 µl PBS. The suspensions were then centrifuged, hydrogen peroxide or PBS was decanted and the spores were washed twice with PBS. Washed spores were plated on YCFA media with 0.1% sodium taurocholate to stimulate spore germination and colony forming units were counted 24 hours later. The experiment was performed independently 3 times for each strain. Clade A strains used were TL178 (RT002/ PG1), TL174 (RT015/ PG1), R20291 (RT027/ PG2), CF5 (RT017/ PG3) and CD305 (RT023/ PG3). Clade B strains used were MON024 (RT033), CDM120 (RT078), WA12 (RT291), WA13 (RT228) and MON013 (RT127). Data were presented using GraphPad Prism v7.03.

***In vivo C. difficile* colonization experiment**

Five female 8-week-old C57BL/6 mice were given 250 mg/l clindamycin (Apollo Scientific) in drinking water. After 5 days, clindamycin treatment was interrupted and 100 mM of glucose, fructose or ribose was added to mouse drinking water for the rest of the experiment; no sugars were given to control mice. After 3 days, mice were infected orally

with 6×10^3 spore/mouse of *C. difficile* R20291 (RT027) or M120 (RT078) strain. Fecal samples were collected from all mice before infection to check for pre-existing *C. difficile* contamination. Spore suspensions were prepared as described above¹⁸. After 16 hours, fecal samples were collected from all mice to determine viable *C. difficile* cell counts by serial dilution and plating on CCEY agar supplemented with 0.1% sodium taurocholate. The mean values of 5 mice are presented from 1 representative experiment which was repeated once with similar results. Data were presented using GraphPad Prism version 7.03. Ethical approval for mouse experiments was obtained from the Wellcome Sanger Institute.

Reporting Summary

Further information on research design is available in the Life Sciences Reporting Summary linked to this article.

Data Availability

Genomes have been deposited in the European Nucleotide Archive. Accession codes are listed in Supplementary Table 1. The 13 *C. difficile* reference isolates (Supplementary Table 2) are publicly available from the National Collection of Type Cultures (NCTC) and the annotation of these genomes are available from the Host-Microbiota Interactions Lab (HMIL; www.lawleylab.com), Wellcome Sanger Institute.

Code Availability

No custom code was used.

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638 **Methods-only References**

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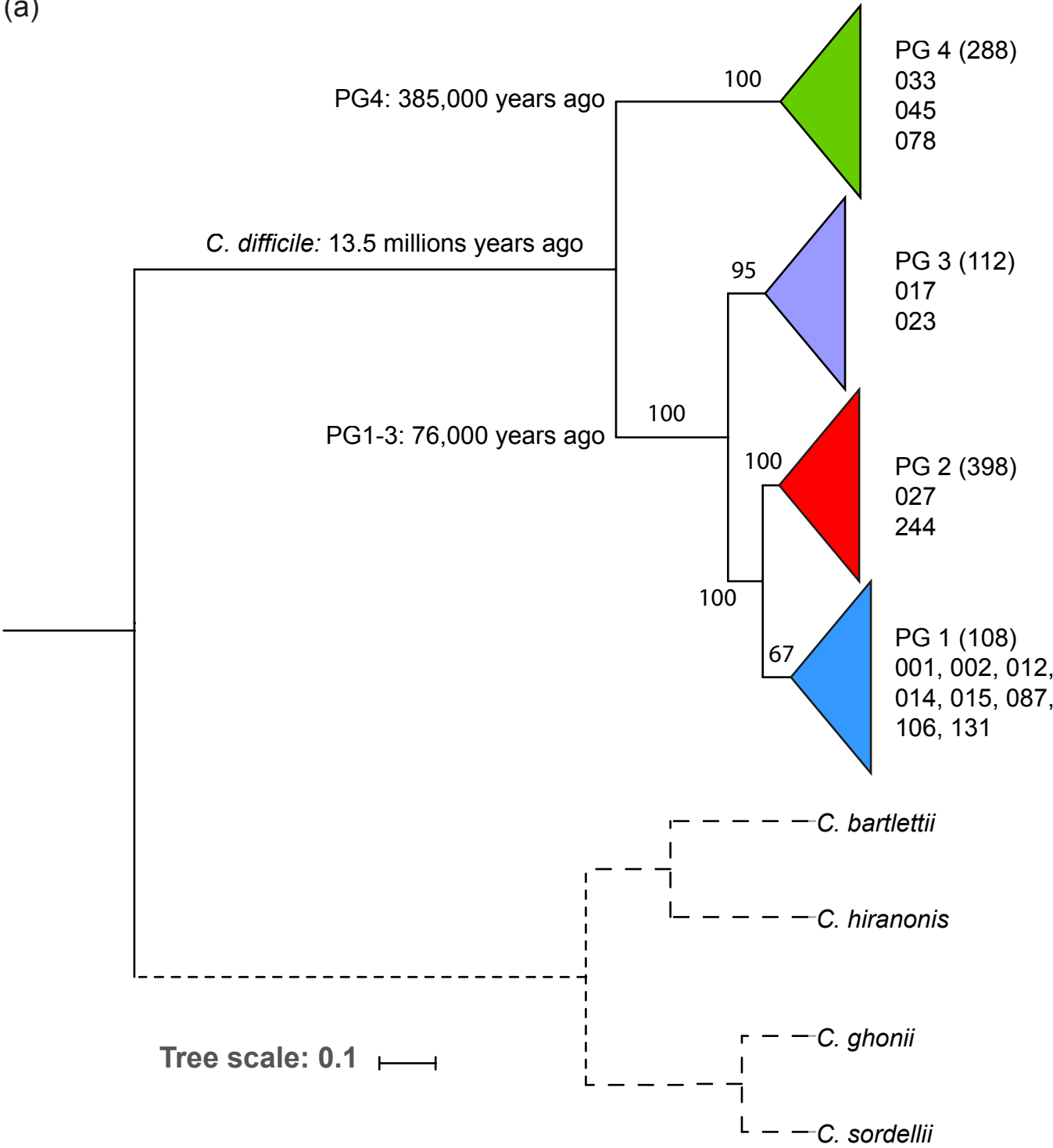
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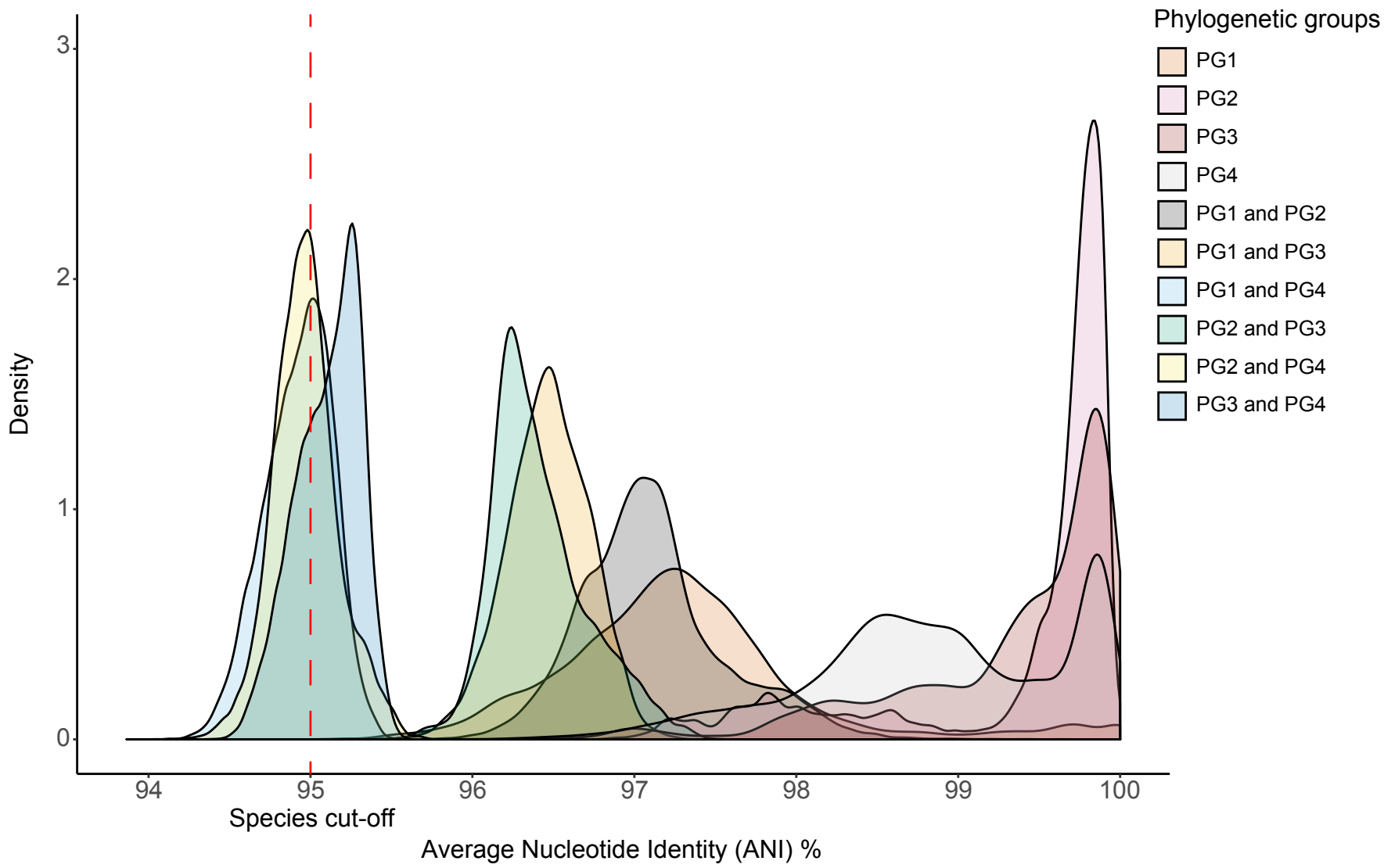
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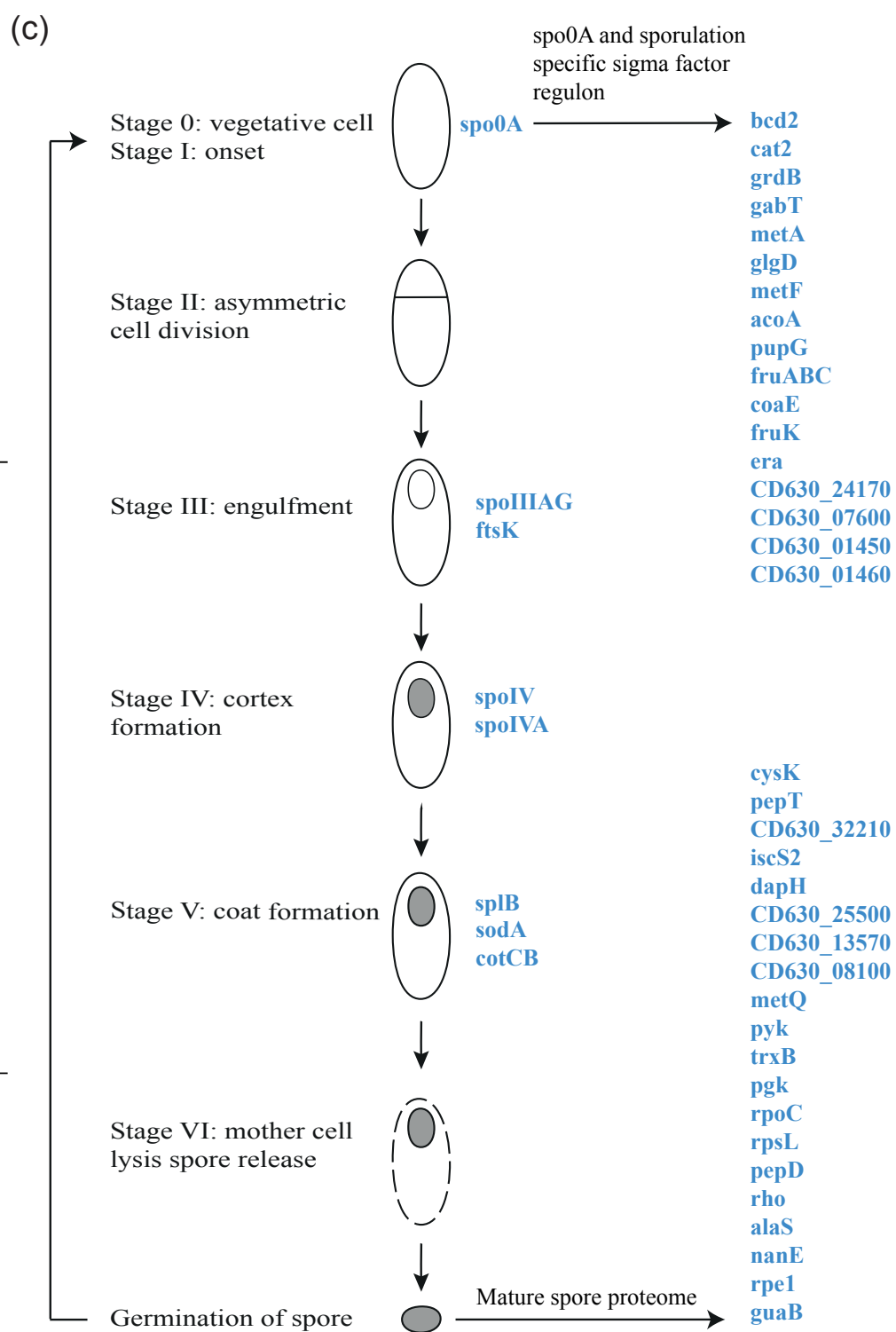
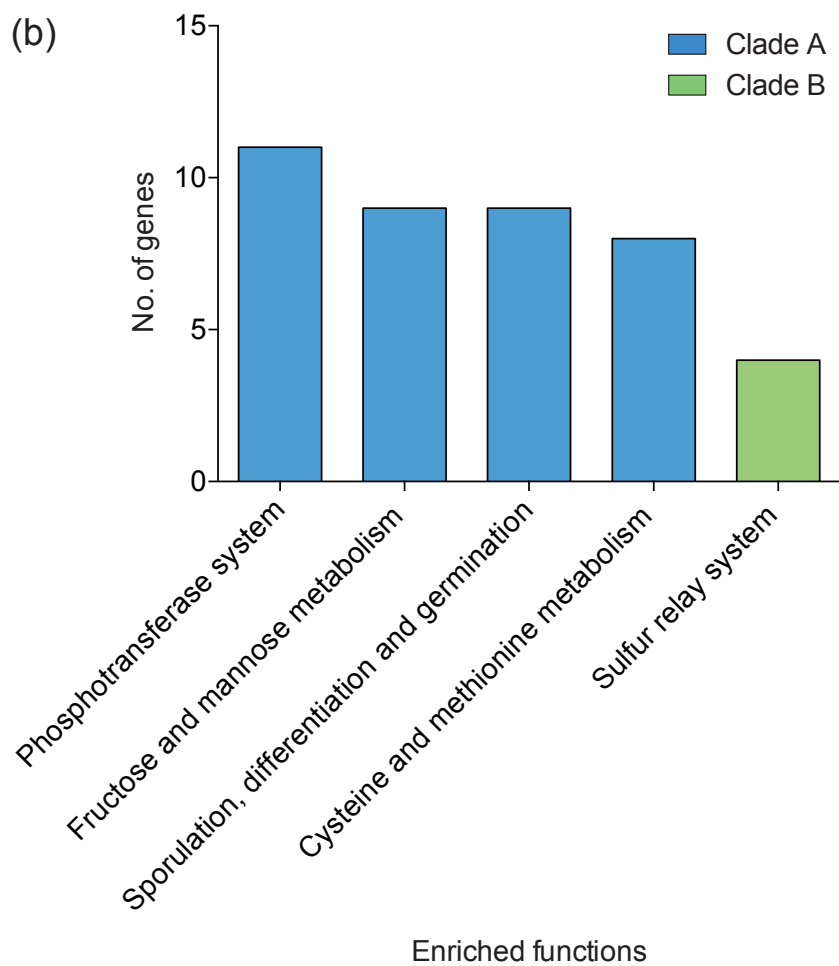
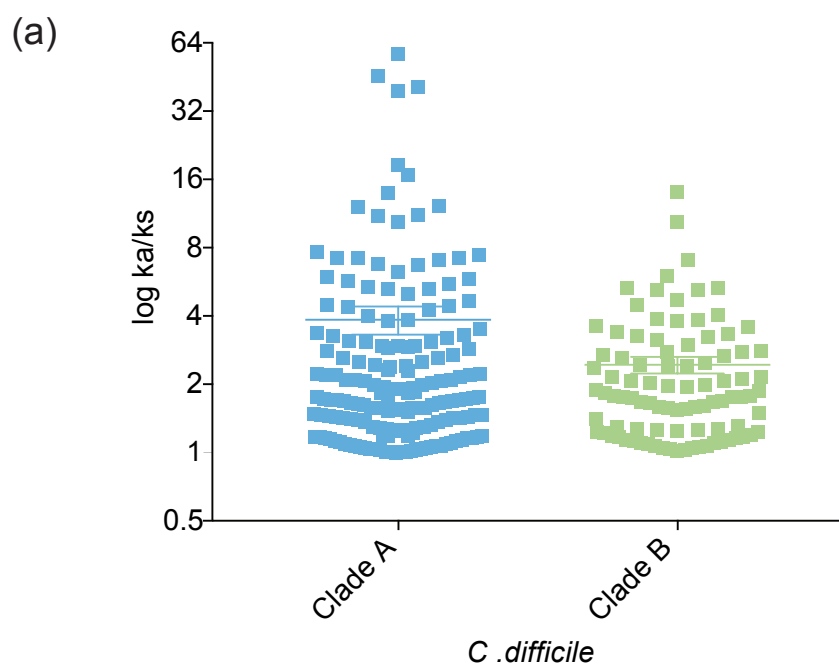
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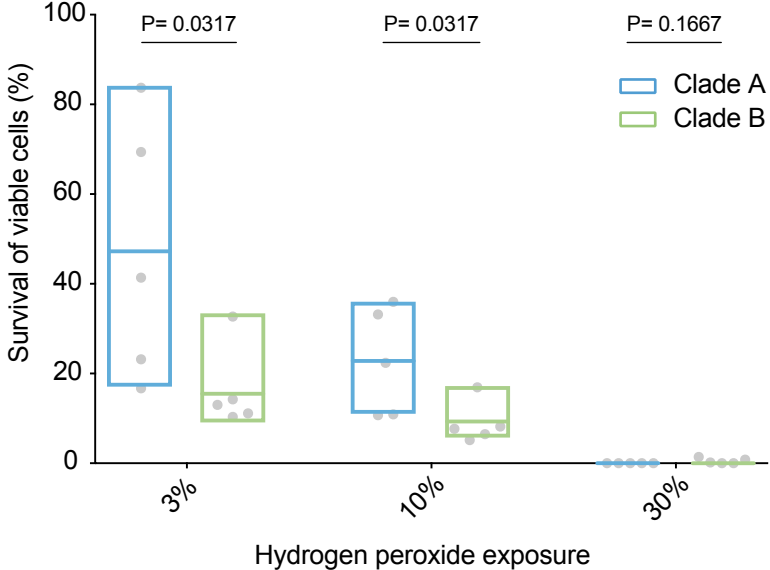


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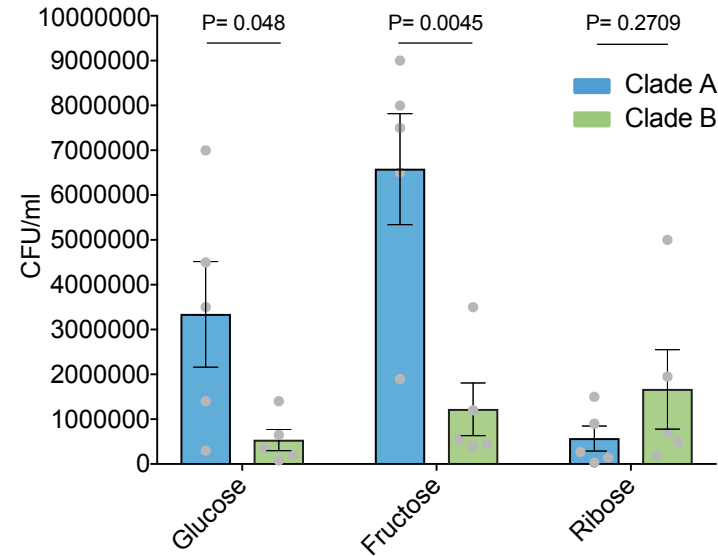




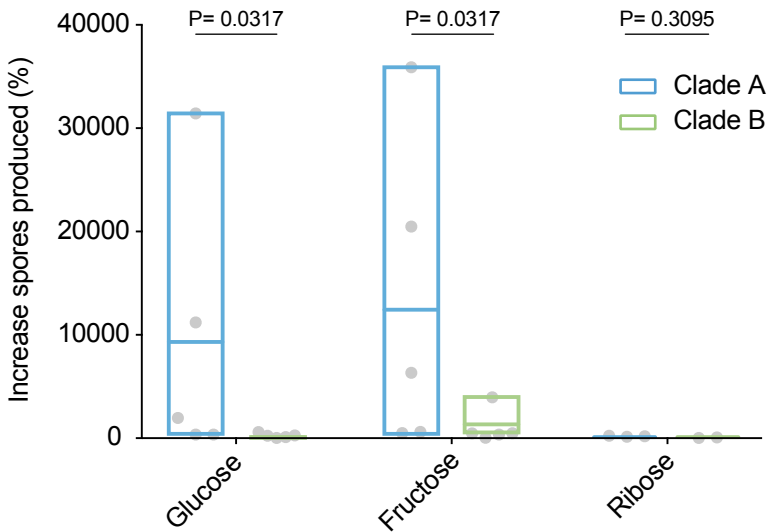
(a) Environmental survival



(b) Host Colonization



(c) Sporulation



(d) Phenotypic adaptations in *C. difficile* Clade A that enhance human transmission

